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LISTING OF THE CLAIMS

1. (Original) A solid formulation comprising at least one antibody, and histidine in a sufficient amount to stabilize said at least one antibody in said solid formulation.

- 2. (Original) The solid formulation of Claim 1, further comprising an excipient.
- 3. (Original) The solid formulation of Claim 2, wherein said at least one other excipient is selected from the group consisting of: mannitol, Polysorbate 20, Polysorbate 80, succinate, citrate, Tris, phosphate, sucrose, trehalose, amino acids, polyols, PEG, BSA, sucrose, lactose, maltose, and sorbital.
- 4. (Original) The solid formulation of Claim 2, wherein said at least one other excipient is arginine.
- 5. (Original) The solid formulation of Claim 1, wherein said at least one antibody is a mammalian antibody.
- 6. (Original) The solid formulation of Claim 1, wherein said at least one antibody is a human antibody.
- 7. (Original) The solid formulation of Claim 1, wherein said at least one antibody is a human monoclonal IgG₂ antibody.
- 8. (Original) The formulation of Claim 1, wherein the sufficient amount of histidine is between 6 and 40 mM.
- 9. (Original) The formulation of Claim 1, wherein the sufficient amount of histidine is about 15 mM of histidine.
 - 10. (Original) A method of preparing an antibody in a solid formulation comprising: mixing at least one antibody with a stabilizing amount of histidine to form a mixture; and

treating said mixture to generate a solid formulation of said antibody and said histidine.

- 11. (Original) The method of Claim 10, wherein treating said mixture comprises lyophilizing said mixture.
- 12. (Original) The method of Claim 10, wherein said solid formulation is a lyophilized cake.
 - 13. (Original) The method of Claim 11, wherein lyophilizing said mixture comprises:

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freezing said mixture at a rate of about -0.35° C per minute until said mixture reaches a temperature of about -45° C; and

sufficiently drying said mixture.

- 14. (Original) The method of Claim 13, wherein drying comprises a primary and a secondary drying.
- 15. (Original) The method of Claim 12, further comprising reconstituting said lyophilized cake with a reconstituting agent.
- 16. (Original) The method of Claim 15, wherein said reconstituting agent comprises water for injection (WFI).
- 17. (Original) The method of Claim 10, further comprising adding at least one other excipient to said mixture.
- 18. (Original) The method of Claim 17, wherein said at least one other excipient is selected from the group consisting of: mannitol, Polysorbate 20, Polysorbate 80, succinate, citrate, Tris, phosphate, trehalose, amino acids, polyols, PEG, BSA, sucrose, lactose, maltose, and sorbital.
- 19. (Original) The method of Claim 15, wherein said at least one other excipient is arginine.
- 20. (Original) The method of Claim 10, wherein the stabilizing amount of histidine is between 6-40 mM.
- 21. (Original) The method of Claim 10, wherein the stabilizing amount of histidine is about 15 mM.
- 22. (Original) The method of Claim 11, wherein lyophilizing said mixture occurs in less than 100 hours.
- 23. (Original) The method of Claim 11, wherein lyophilizing said mixture occurs in less than 50 hours.
- 24. (Original) The method of Claim 11, wherein lyophilizing said mixture occurs in about 45 hours.
- 25. (Original) A kit for preparing a solid formulation of a stabilized antibody comprising;

a first container, comprising at least one antibody in solution, and

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a second container comprising a sufficient amount of histidine in solution to stabilize said antibody when said antibody is dried into a solid formulation.

- 26. (Original) The kit of Claim 25, wherein said antibody is a mammalian antibody.
- 27. (Original) The kit of Claim 25, wherein said antibody is a human antibody.
- 28. (Original) The kit of Claim 25, wherein said antibody is a human monoclonal IgG₂ antibody.
- 29. (Original) The kit of Claim 25, wherein the sufficient amount of histidine is between 6-40 mM.
- 30. (Original) The kit of Claim 25, wherein the sufficient amount of histidine is about 15 mM.
- 31. (Original) A liquid formulation comprising at least one antibody, and histidine in a sufficient amount to stabilize said at least one antibody in said liquid formulation.
 - 32. (Original) The liquid formulation of Claim 31, further comprising an excipient.
- 33. (Original) The liquid formulation of Claim 32, wherein said at least one other excipient is selected from the group consisting of: mannitol, Polysorbate 20, Polysorbate 80, succinate, citrate, Tris, phosphate, sucrose, trehalose, amino acids, polyols, PEG, BSA, sucrose, lactose, maltose, and sorbital.
- 34. (Original) The liquid formulation of Claim 32, wherein said at least one other excipient is arginine.
- 35. (Original) The liquid formulation of Claim 31, wherein said at least one antibody is a mammalian antibody.
- 36. (Original) The liquid formulation of Claim 31, wherein said at least one antibody is a human antibody.
- 37. (Original) The liquid formulation of Claim 31, wherein said at least one antibody is a human monoclonal IgG_2 antibody.
- 38. (Original) The liquid formulation of Claim 31, wherein the sufficient amount of histidine is between 6 and 40 mM.
- 39. (Original) The formulation of Claim 31, wherein the sufficient amount of histidine is about 15 mM of histidine.